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Reduction of imines with a reusable bimetallic PdCo–Fe₃O₄ catalyst at room temperature under atmospheric pressure of H₂†

Sabyuk Yang and Byeong Moon Kim *

Bimetallic nanocatalysts have been used for the development of organic reactions, owing to the synergistic effect between the transition metals. A new procedure for synthesizing amines by the reduction of imines with H₂ at atmospheric pressure and room temperature in the presence of PdCo–Fe₃O₄ nanoparticles is reported. The straightforward procedure, mild reaction conditions, high turnover number, and recyclability extend the scope of this reaction to practical applications.

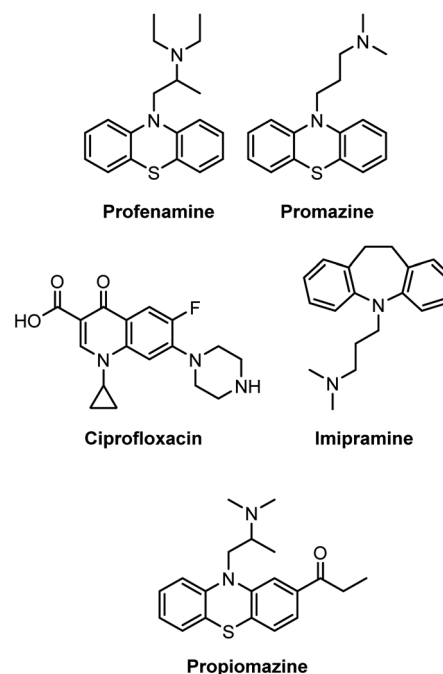
Introduction

Amines are versatile organic molecules found in various natural products,¹ drug molecules,² and dyes.³ Of particular interest is a large number of pharmaceuticals, such as profenamine (antidyskinetic),⁴ promazine (antiemetic),⁵ ciprofloxacin (antibiotic),⁶ imipramine (antidepressant),⁷ and propiomazine (antipsychotic agent),⁸ which all contain the amine functionality (Scheme 1). In this context, research to develop continually mild and convenient synthetic procedures for synthesizing amines from various precursors remains highly relevant to the present pharmaceutical industry.⁹

Among many established procedures, the reduction of imines is one of the most straightforward methods, owing to the fact that imines can be readily produced from precursor amines and carbonyl compounds.¹⁰ Although the reduction of imines can be facilitated by adding stoichiometric hydride reagents, such as NaBH₄,¹¹ NaBH₃CN,¹² or NaBH(OAc)₃,¹³ catalytic processes are preferred for a large-scale production. Various homogeneous transition metal catalysts such as Pd,^{14a} Ir,^{15a} Ru,^{16a,b} Mn,^{17a} and Mo¹⁸ have been employed in the reduction of imines using H₂. Drawbacks associated with reactions on these catalysts include high temperature and high H₂ pressure required to achieve high turnover numbers (TON). In the meantime, heterogeneous catalysis employing transition metals have been gaining increasing attention as an alternative to homogeneous catalysis, owing to the usability and recyclability.¹⁹ For the purpose of overcoming the shortcomings mentioned above, efforts have been concentrated on recycling the catalyst, thereby allowing repeated applications after the hydrogenation process.^{14b–e,15b,c,16c,17b} Therefore, we

envisioned that further development of the catalytic reaction reduction of imines under mild reaction conditions, *i.e.*, room temperature and at atmospheric pressure of H₂, with a high turnover number, and good catalyst reusability is still in need (Table 1).

There has been increasing interest in bimetallic heterogeneous catalysts because they often exhibit a synergistic effect between two types of metals, which is not observed in metals of single type.²⁰ A series of bimetallic heterogeneous catalysts, supported on iron oxide (Fe₃O₄) nanoparticles (NPs), have recently been developed for application in the development of



Scheme 1 Amines in some commercially available drugs.

Department of Chemistry, College of Natural Sciences, Seoul National University, Seoul 08826, South Korea. E-mail: kimbm@snu.ac.kr

† Electronic supplementary information (ESI) available: Catalyst characterization, experimental details, and ¹H and ¹³C NMR spectra. See DOI: 10.1039/d1ra08552b

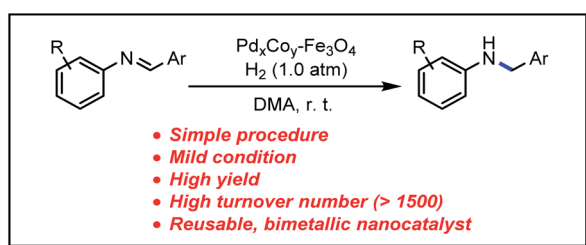


Table 1 Representative reports on the reduction of imines with various transition metal catalysts in the presence of H₂

Entry	Metal	Condition	TON	Note	Ref.
1	Pd	r. t., 40 atm H ₂	20	HM ^a	14a
2	Ir	80 °C, 120 atm H ₂	3960	HM	15a
3	Ru	40 °C, 50 atm H ₂	31 500	HM	16a
4	Ru	70 °C, 5.0 atm H ₂	1000	HM	16b
5	Mn	50 °C, 50 atm H ₂	49	HM	17a
6	Mo	r. t., 30 atm H ₂	333	HM	18a
7	Mo	140 °C, 60 atm H ₂	800	HM	18b
8	Pd	r. t., 1.0 atm H ₂	404	HT ^b	14c
9	Pd	r. t., 1.0 atm H ₂	8	HT	14e
10	Ir	50 °C, 100 atm H ₂	12 400	HT	15b
11	Ir	r. t., 10 atm H ₂	1188	HT	15c
12	Ru	90 °C, 30 atm H ₂	1	HT	16c
13	Mn	r. t., 5.0 atm H ₂	19	HT	17b
14	PdCo	r. t., 1.0 atm H ₂	1583	HT	This work

^a HM = homogeneous catalysis. ^b HT = heterogeneous catalysis.

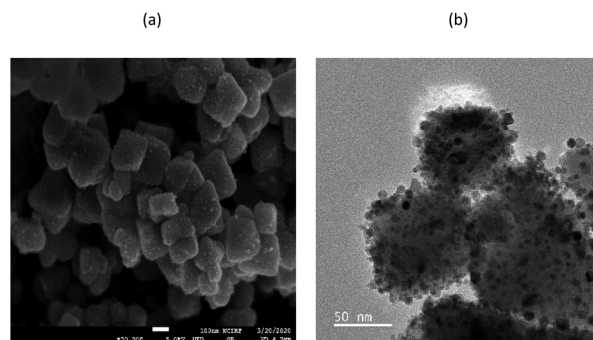
various organic reactions. This included the development of PdPt-Fe₃O₄ NPs that have been employed in nitroarene reduction,²¹ arylsilylation,²² as well as secondary amine synthesis from benzonitriles and nitroarenes.²³ Additionally, AuPd-Fe₃O₄ NP catalysts have been developed for reductive amination,²⁴ oxidation of furan-2,5-dimethylcarboxylate,²⁵ and N-formylation of secondary amines.²⁶ However, all these catalysts employed expensive, precious metals (Pd and Pt, Au and Pd) among the transition metal element group. Further research therefore focused on the development of a bimetallic nanocatalyst that incorporated at least one inexpensive transition metal. Accordingly, PdCo bimetallic catalysts were identified as a promising alternative, with applications in various organic transformations.²⁷ These include the development of a PdCo catalyst for the transfer hydrogenation of carbonyl compounds by Kumar *et al.*,^{27a} while Wang group employed a PdCo electrocatalyst for the reduction of oxygen and the oxidation of ethanol.^{27b} Chen group also successfully employed a PdCo alloy for oxygen reduction,^{27c} while Salmeron group investigated the effect of a PdCo nanocatalyst in CO oxidation.^{27d} A PdCo bimetallic catalyst has, however, not yet been employed in imine reduction. In the present study, we have developed PdCo-Fe₃O₄ as a recyclable bimetallic nanocatalyst for the reduction of imines (Scheme 2) under very mild conditions.

**Scheme 2** Synthesis of amines from imines using reusable PdCo-Fe₃O₄ bimetallic catalyst under mild condition.

Results and discussion

Catalyst characterization

Here, bimetallic PdCo-Fe₃O₄ NPs were synthesized through the modification of an existing synthetic procedure for AuPd-Fe₃O₄.²⁴ Sequentially adding palladium(II) chloride (PdCl₂) in ethylene glycol (EG), cobalt(II) chloride hexahydrate (CoCl₂·6H₂O) in water, and aqueous sodium borohydride solution dropwise to a Fe₃O₄ solution in water followed by stirring under 60 °C for 24 h afforded PdCo-Fe₃O₄ NPs. The synthesized nanocatalyst was characterized employing various techniques. Scanning electron microscopy (SEM) images (Fig. 1a and S1†) were obtained using a JSM-7800F Prime microscope (JEOL Ltd., Tokyo, Japan). High-resolution transmission electron microscopy (HR-TEM) images (Fig. 1b, S4 and S6†) were obtained using a JEM-3010 microscope (JEOL Ltd., Tokyo, Japan), indicating successful deposition of Pd and Co atoms on the Fe₃O₄ support. Scanning transmission electron microscopy images (STEM, Fig. 2 and S5†) were obtained using a JEM-ARM200F microscope (JEOL Ltd., Tokyo, Japan), while energy-dispersive X-ray spectroscopy (EDS) images (Fig. S2 and S3†) were obtained using a JSM-7800F Prime Field Emission Scanning Electron Microscope (JEOL Ltd., Tokyo, Japan), showing that Pd and Co were randomly distributed. The electron energy loss spectroscopy (EELS) was performed using a Themis Z STEM (ThermoFisher, MA, USA), where the high-energy loss spectrum images indicated that the Co lines are located at 780–800 eV (Fig. S7†). In addition, the X-ray diffraction (XRD) data obtained using a D8 Advance XRD (Bruker, Billerica, MA, USA) indicated that the diffraction peaks at ~40° and 46–47° 2θ were shifted. This suggested that PdCo-Fe₃O₄ is made up with Pd-Co bimetallic alloy (Fig. S8†).^{27h} X-ray photoelectron spectroscopy (XPS) data were obtained using a SIGMA PROBE XPS (ThermoFisher Scientific, UK) and provided information on the electronic states of the catalyst (Fig. S9†). Here, first major peaks at the Pd 3d_{5/2} and Pd 3d_{3/2} energy levels were observed at 335.35 and 340.61 eV, respectively, which correspond to metallic Pd(0) species.²⁸ In addition, first major peaks at the Co 2p_{3/2} and Co 2p_{1/2} energy levels were observed at 779.84 and 795.64 eV, respectively, corresponding to metallic Co(0) species.²⁹ Finally, an OPTIMA 8300 inductively coupled plasma-atomic emission spectrometer (ICP-AES) (PerkinElmer, Waltham, MA, USA), was

**Fig. 1** (a) SEM and (b) HR-TEM image of PdCo-Fe₃O₄ catalyst.

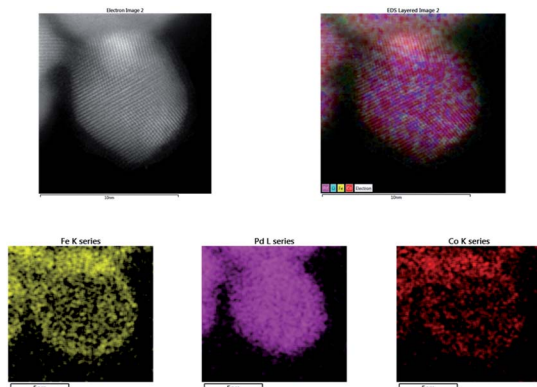


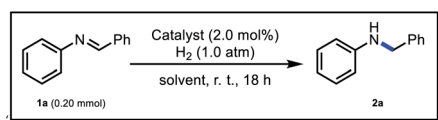
Fig. 2 STEM-EDS image of PdCo-Fe₃O₄ catalyst.

employed to determine that the PdCo-Fe₃O₄ NPs consisted of 7.49 wt% Pd and 3.89 wt% Co, at a molar ratio of 0.99 : 1.00 (Fig. S11[†]). The synthesized PdCo-Fe₃O₄ NPs were subsequently utilized to optimize the imine reduction reaction.

Reaction optimization

For the initial screening study, *N*-benzylideneaniline (**1a**) was selected as a representative substrate and *N,N*-dimethylacetamide (DMA) as the solvent. Various reaction conditions were examined, with the results presented in Tables 2 and S1–S4.[†]

Table 2 Screening data of the synthesis of amine from imine reduction^a



Entry	Catalyst	Reductant	Solvent	Yield ^b	TON ^c
1	Fe ₃ O ₄	H ₂	DMA	N. D. ^d	—
2 ^e	Pd/C	H ₂	DMA	N. D.	—
3 ^f	Pd-Fe ₃ O ₄	H ₂	DMA	60	15
4 ^g	Co-Fe ₃ O ₄	H ₂	DMA	N. D.	—
5	PdCo-Fe ₃ O ₄	H ₂	DMA	95 (91 ^h)	24
6	PdCo-Fe ₃ O ₄	H ₂	MeOH	71	18
7	PdCo-Fe ₃ O ₄	H ₂	DMF	83	21
8 ⁱ	PdCo-Fe ₃ O ₄	PhSiH ₃	DMA	2	0.50
9 ⁱ	PdCo-Fe ₃ O ₄	BH ₃ NH ₃	DMA	47	12
10 ⁱ	PdCo-Fe ₃ O ₄	NaBH ₄	DMA	39	10
11 ^j	PdCo-Fe ₃ O ₄	H ₂	DMA	>99	50
12 ^k	PdCo-Fe ₃ O ₄	H ₂	DMA	98 (97 ^h)	98
13 ^l	PdCo-Fe ₃ O ₄	H ₂	DMA	95	1583

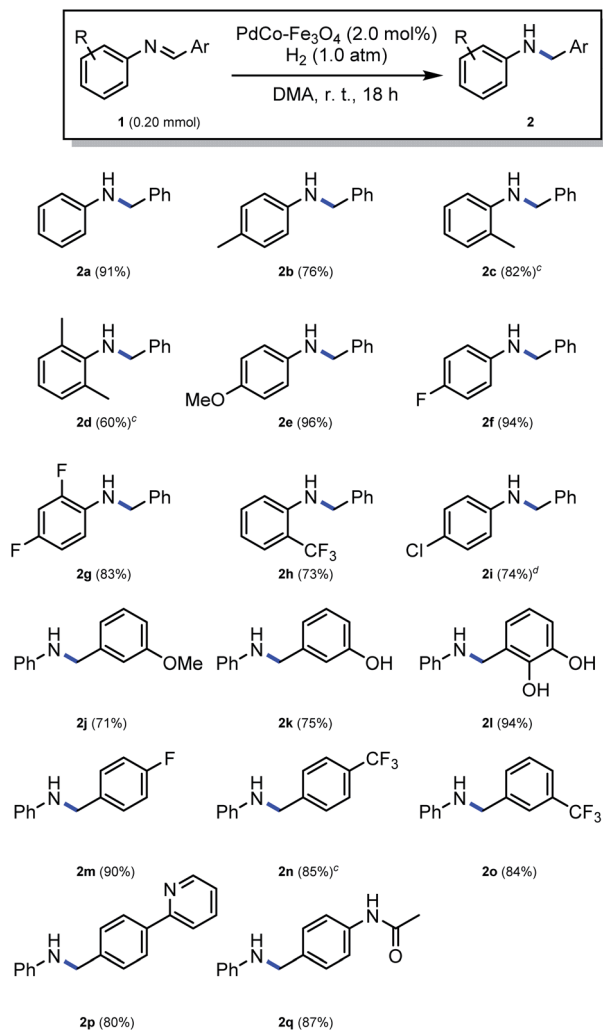
^a Reaction conditions: **1a** (0.20 mmol), catalyst (2.0 mol%), H₂ (1.0 atm), solvent (1.0 mL), r. t., 18 h. ^b Determined from ¹H NMR spectral analysis through the use of anisole as an internal standard. ^c Turnover number (TON) = mmol of product/mmol of total metal except Fe. ^d N. D. = not detected. ^e Pd/C (4.0 mol%) was used as a catalyst. ^f Pd-Fe₃O₄ (4.0 mol%) was used as a catalyst. ^g Co-Fe₃O₄ (4.0 mol%) was used as a catalyst. ^h Yield of isolated product. ⁱ Result with 3.0 equiv. of reductant instead of 1.0 atm of H₂. ^j PdCo-Fe₃O₄ (1.0 mol%) was used as a catalyst. ^k PdCo-Fe₃O₄ (0.50 mol%) was used as a catalyst. ^l Result with **1a** (3.3 mmol), PdCo-Fe₃O₄ (0.03 mol%), 78 h.

When the reaction was carried out with only Fe₃O₄ as a catalyst under 1.0 atm of H₂, the reaction did not proceed and **1a** remained unchanged (entry 1). Although the reaction resulted in hydrogenolysis side products when Pd/C was employed as a catalyst, none of the desired product was produced (entry 2). However, employing 4.0 mol% of Pd-Fe₃O₄ resulted in a 60% yield of the desired product **2a**, while catalysis employing Co-Fe₃O₄ alone did not produce any **2a** (entries 3 and 4, respectively). Finally, employing the PdCo-Fe₃O₄ bimetallic catalyst resulted in a 95% yield, indicating an outstanding synergistic effect between Pd and Co (entry 5). To confirm this synergy, the reaction yields of several substrates from the reactions employing monometallic catalysts were compared (Table S6[†]). Here, the reaction yields using PdCo-Fe₃O₄ were higher than those obtained when Pd-Fe₃O₄ or Co-Fe₃O₄ was used independently. The initial kinetics associated with employing either monometallic catalysts or PdCo-Fe₃O₄ catalyst were also investigated (Fig. S17[†]). An enhanced reaction rate was observed for the reaction when the PdCo-Fe₃O₄ catalyst was employed, demonstrating an additional benefit associated with the use of the bimetallic catalyst. The effect of other solvents was also evaluated. Here, employing methanol and *N,N*-dimethylformamide (DMF) (entries 6 and 7), resulted in lower yields when compared to yields obtained with DMA. Reactions with other reducing agents such as phenylsilane (entry 8), ammonia borane (entry 9), and sodium borohydride (entry 10) also resulted in lower product yields. Ultimately, approximately quantitative yields were obtained (entries 11 and 12, respectively) even when the catalyst loading was reduced from 2.0 mol% to 1.0 or 0.50 mol%, respectively. Furthermore, the turnover number (TON) of the catalyst could be higher than 1500 even when the catalyst loading was low, demonstrating the efficiency of the catalyst to facilitate imine reduction (entry 13).

Substrate scope

The reduction of several imines with various substituents on the aromatic ring was subsequently evaluated at the optimized conditions, employing a 0.20 M substrate concentration for a period of 18 h. Here, as shown in Scheme 3, the reduction of *N*-benzylideneaniline, with no substituent on the aniline or benzene ring, proceeded smoothly, and a 91% yield of the desired product (**2a**) could be isolated. Substrates with a methyl group on the aniline moiety resulted in moderate to good yields (76% and 82% for **2b** and **2c**, respectively), while the reaction of a substrate with a single methyl group at position 2 of the aromatic ring did not have an impact on the yield (**2c**). However, substitution with two methyl groups at positions 2 and 6 resulted in a low yield (**2d**), demonstrating the influence of steric hindrance. Strong electron-donating methoxy group on the aromatic ring was well tolerated (**2e**). The reduction of substrates with strong electron-withdrawing groups, such as fluorine, were associated with high yields (94% and 83% for **2f** and **2g**, respectively). The reduction of an imine containing a relatively bulky trifluoromethyl group at position 2 of the aniline ring provided a 73% yield (**2h**). For the reduction of substrates containing a halogen such as Cl, Br, and I, the use of





Scheme 3 Substrate scope of the synthesis of amines by the reduction of imines.^{a,b} ^aResult with **1** (0.20 mmol), PdCo-Fe₃O₄ (2.0 mol%), H₂ (1.0 atm), DMA (1.0 mL), r. t., 18 h. ^bYield of isolated product. ^cResult with **1** (0.50 mmol) in DMA (2.0 mL). ^dPd_{0.05}Co₁-Fe₃O₄ (0.10 mol%), r. t., 30 h.

PdCo-Fe₃O₄ was not appropriate, because substrate dehalogenation occurred.³⁰ It was therefore hypothesized that reducing the content of Pd relative to that of Co to a minimum, may suppress dehalogenation while maintaining the synergy between Pd and Co. This was confirmed as the reaction with Pd_{0.05}Co₁-Fe₃O₄ proceeded successfully without knocking out the chloride group at position 4 of the aniline ring, resulting in a 74% yield obtained within 30 h (**2i**). Various substrates with substituents on the benzyl ring were further tested. Here, the reaction of a substrate containing a methoxy group at position 3 provided a 71% yield (**2j**), while a good yield (75%, **2k**) was also obtained using a 3-hydroxy substituted substrate. The reaction of a substrate with multiple hydroxyl groups at positions 2 and 3 on the benzyl ring also resulted in a high yield (94%, **2l**). Similarly, when a substrate possessing fluoro or trifluoromethyl group on the benzyl ring was employed in the reaction, high yields (90%, 85% and 84% for **2m**, **2n** and **2o**, respectively) could

be obtained under similar conditions. Reduction of a substrate containing a heteroaromatic ring, such as pyridine was also well tolerated, resulting in an 80% yield (**2p**). Finally, an 87% yield was obtained when a substrate with an acetamide group on the benzyl ring was employed in the reaction. In this instance, the amide functionality was intact (**2q**).

Effect of Pd : Co ratio and various supports

Based on the results from Table 2, the role of Co in the PdCo bimetallic catalyst appeared to accelerate hydrogenation since the bimetallic catalyst works more efficiently than either of the monometallic catalyst. Therefore, we were curious whether any differences in catalytic activity could be attributed to the composition of the two metals in the catalyst. Catalysts with various Pd : Co ratios were therefore prepared (Fig. S12†) and tested under the same reaction conditions (Table 3, entries 1–6). Here, the experimental data indicated that the yields were excellent regardless of the Pd : Co molar ratios. Reaction using the catalyst with 0.04 : 1 molar ratio of Pd : Co resulted in 91% yield in 18 h (Table 3, entry 1). In an attempt to evaluate the effect of the Fe₃O₄ support on the catalyst performance, PdCo catalysts with various supports were synthesized (Fig. S16 and Table S5†) and the reactions employing catalysts with new supports were tested (Table 3, entries 7–9). Although PdCo-CeO₂ (entry 8) exhibited a similar reactivity to that of PdCo-Fe₃O₄, it was concluded that PdCo-Fe₃O₄ offered additional advantages owing to the simplicity associated with the recycling process. This will be discussed in the following section.

Recycling test

Considering that heterogeneous catalysts are often employed based on their convenient recovery and the possibility to reuse the catalyst, the recyclability of various Pd_xCo_y-Fe₃O₄ NPs was tested under optimized conditions (Fig. 3 and S13†). PdCo-

Table 3 Reactivity comparison of various PdCo catalysts^a

Entry	Catalyst	Yield ^b
1	Pd _{0.04} Co ₁ -Fe ₃ O ₄	91
2	Pd _{0.26} Co ₁ -Fe ₃ O ₄	96
3	Pd _{0.46} Co ₁ -Fe ₃ O ₄	96
4	Pd ₁ Co ₁ -Fe ₃ O ₄	95
5	Pd ₁ Co _{0.45} -Fe ₃ O ₄	93
6	Pd ₁ Co _{0.28} -Fe ₃ O ₄	93
7	PdCo-TiO ₂	82
8	PdCo-CeO ₂	93
9	PdCo-C	63

^a Reaction conditions: **1a** (0.20 mmol), catalyst (2.0 mol%), H₂ (1.0 atm), DMA (1.0 mL), r. t., 18 h. ^b Determined from ¹H NMR spectral analysis through the use of anisole as an internal standard.



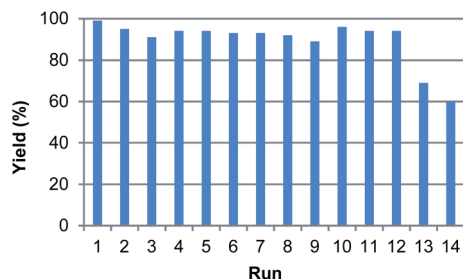
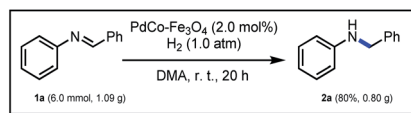


Fig. 3 Results of recycling test under PdCo-Fe₃O₄ NPs.



Scheme 4 A gram scale reaction.

Fe₃O₄ NPs were used as a representative catalyst, after being recovered using a small magnet and washed with MeOH several times. In the recycling experiment with **1a**, a consistent catalytic activity was observed for 12 cycles. However, starting from the 13th cycle, a reduction in reactivity was observed. Evaluation of the SEM and HR-TEM images of the catalyst after the 14th cycle indicated that the used PdCo particles exhibited a high degree of agglomeration as well as detachment of metal atoms from the Fe₃O₄ support (Fig. S14 and S15[†]). Additionally, ICP-AES analysis of the used catalyst showed a reduction in both Pd and Co contents, from 7.49 wt% to 1.69 wt% for Pd and from 3.89 wt% to 0.28 wt% for Co, respectively (Fig. S11[†]), reflecting the decreased reactivity after the repeated catalytic cycles.

Gram scale reaction

The scalability of the reaction (Scheme 4) was further investigated, and at a 6.0 mmol scale, the reaction of **1a** resulted in an 80% yield of **2a**, demonstrating its potential suitability in organic synthesis applications.

Conclusions

In summary, a new catalytic system was developed for the synthesis of amines *via* the reduction of imines. Utilizing a PdCo bimetallic nanocatalyst, various imines could be reduced to amines at room temperature and under 1.0 atm of H₂. This process therefore allowed the catalytic reduction of imines to amines without the use of toxic or polluting stoichiometric reducing agents. Furthermore, the catalytic performance of the bimetallic PdCo-Fe₃O₄ NPs was superior to that of monometallic NPs, such as Pd-Fe₃O₄ and Co-Fe₃O₄. This indicated the existence of a synergistic effect between Pd and Co. The application scope of the reaction was broad, and it could tolerate substrates with various functional groups. The reduction reaction was further shown to be independent of the Pd : Co ratio, while the Fe₃O₄ NPs were proven to be a more effective support material than other supporting materials such

as TiO₂, CeO₂, and C. The PdCo-Fe₃O₄ NPs could be used in more than 10 cycles without a significant loss of catalytic activity. Moreover, it was possible to run the reaction on a gram scale. The high turnover number combined with the simple experimental procedure overcomes the limitations associated with previous approaches such as high reaction temperatures, high H₂ pressures, as well as the use of expensive transition metal precursors. This catalyst will be investigated further to expand its potential applications in organic chemistry.

Experimental

Materials

All commercially available chemicals were purchased from Acros Organics (Pittsburgh, PA, USA), Sigma-Aldrich (St. Louis, MO, USA), Alfa Aesar (Ward Hill, MA, USA), Tokyo Chemical Industry (Tokyo, Japan), and used without further purification. Imine substrates were synthesized by a known procedure.³¹

Synthesis of PdCo-Fe₃O₄ NPs

Initially, 88.7 mg (0.50 mmol) of PdCl₂ and 1.00 g of PVP (*M_w* ~ 10 000 g mol⁻¹, 0.10 mmol) were placed in 20 mL of ethylene glycol (EG) in a 100 mL round-bottom flask. This mixture was sonicated for 10 min and stirred for 1 h at 100 °C in an oil bath. In a separate 100 mL round-bottom flask, 119.0 mg (0.50 mmol) of CoCl₂·6H₂O and 500 mg of PVP (0.050 mmol) were added to 20 mL of water. This mixture was sonicated for 10 min and stirred for 30 min at 60 °C in an oil bath. Meanwhile, 500 mg of Fe₃O₄ NPs were added to 150 mL of water in a two-necked 500 mL round-bottom flask and then sonicated for 10 min. The prepared Pd precursor solution was then injected dropwise onto the Fe₃O₄ suspension with vigorous stirring. After 5 min, the Co precursor solution was added and 90 mg (2.38 mmol) of sodium borohydride in 20 mL of water was injected dropwise. The resulting mixture was stirred for 24 h at 60 °C. Subsequently, the PdCo alloy on Fe₃O₄ nanoparticles was retrieved *via* sonication and washing with ethanol (40 mL × 10 times) and dried on a rotary evaporator to give PdCo-Fe₃O₄ NPs (550 mg, 77% yield based on PdCl₂).

Synthesis of Pd-Fe₃O₄ NPs

Initially, 177 mg of PdCl₂ (1.0 mmol) and 2.00 g of PVP (0.20 mmol) were placed in 40 mL of EG in a 100 mL round-bottom flask. This mixture was sonicated for 10 min and stirred for 1 h at 100 °C. Meanwhile, 500 mg of Fe₃O₄ NPs was added to 150 mL of EG in a two-necked 500 mL round-bottom flask. The prepared precursor solution was then injected dropwise to Fe₃O₄ NPs in 150 mL of EG, and stirred at 100 °C for an additional 24 h. The resultant product was washed with ethanol (40 mL × 10 times) and dried on a rotary evaporator to give Pd-Fe₃O₄ NPs (440.0 mg, 36% yield based on PdCl₂).

Synthesis of Co-Fe₃O₄ NPs

Initially, 23.8 mg of CoCl₂·6H₂O (0.30 mmol) and 200 mg of PVP (0.020 mmol) were placed in 4.0 mL of water in a 10 mL round-



bottom flask. This solution was sonicated for 1 min and stirred for 30 min at 60 °C. Meanwhile, 100 mg of Fe₃O₄ NPs was added to 30 mL of water in a two-necked 100 mL round-bottom flask. The prepared precursor solution was then injected dropwise to Fe₃O₄ NPs in 30 mL of water, followed by dropwise addition of 30 mg of sodium borohydride (0.79 mmol) in 4.0 mL of water. The mixture was stirred at 60 °C for an additional 24 h. The resultant product was washed with ethanol (40 mL × 10 times) and dried on a rotary evaporator to give Co-Fe₃O₄ NPs (98.5 mg, 30% yield based on CoCl₂·6H₂O).

Synthesis of Pd_xCo_y-Fe₃O₄ NPs

For Pd_xCo_y-Fe₃O₄ synthesis, the same method used for the synthesis of PdCo-Fe₃O₄ NPs was employed, with different quantities of metals. To prepare Pd_{0.26}Co₁-Fe₃O₄ NPs, PdCl₂ (6.6 mg, 0.038 mmol) with PVP (0.0060 mmol), CoCl₂·6H₂O (35.7 mg, 0.15 mmol) with PVP (0.010 mmol), and sodium borohydride (0.79 mmol) were used. For the synthesis of Pd_{0.46}Co₁-Fe₃O₄ NPs, PdCl₂ (11.1 g, 0.063 mmol) with PVP (0.010 mmol), CoCl₂·6H₂O (35.7 mg, 0.15 mmol) together with PVP (0.010 mmol), and sodium borohydride (0.79 mmol) were used. In the case of Pd₁Pd_{0.45}-Fe₃O₄ NPs, PdCl₂ (17.7 mg, 0.10 mmol) with PVP (0.020 mmol), CoCl₂·6H₂O (18.2 mg, 0.077 mmol) with PVP (0.0050 mmol), and sodium borohydride (0.79 mmol) were used. Finally, for the preparation of Pd₁Co_{0.28}-Fe₃O₄ NPs, PdCl₂ (17.7 mg, 0.10 mmol) with PVP (0.020 mmol), CoCl₂·6H₂O (11.1 mg, 0.047 mmol) with PVP (0.0030 mmol), and sodium borohydride (0.79 mmol) were used. Meanwhile, 100 mg of Fe₃O₄ NPs were added to 30 mL of water in a two-necked 100 mL round-bottom flask and then sonicated for 10 min. The prepared Pd precursor solution in 4.0 mL of EG was then injected dropwise onto the Fe₃O₄ suspension with vigorous stirring. After 5 min, the Co precursor solution in 4.0 mL of water was added and sodium borohydride in 4.0 mL of water was injected dropwise. The resulting mixture was stirred for 24 h at 60 °C. Subsequently, the Pd_xCo_y alloy on Fe₃O₄ nanoparticles was retrieved *via* sonication and washing with ethanol (40 mL × 10 times) and dried on a rotary evaporator to give Pd_{0.26}Co₁-Fe₃O₄ NPs (99.0 mg, 59% yield based on PdCl₂), Pd_{0.46}Co₁-Fe₃O₄ NPs (96.0 mg, 57% yield based on PdCl₂), Pd₁Pd_{0.45}-Fe₃O₄ NPs (98.0 mg, 42% yield based on CoCl₂·6H₂O), and Pd₁Co_{0.28}-Fe₃O₄ NPs (98.0 mg, 69% yield based on CoCl₂·6H₂O).

Catalyst characterization

SEM images were obtained using JSM-7800F Prime (JEOL Ltd., Tokyo, Japan) and MERLIN Compact (ZEISS, Oberkochen, Germany). HR-TEM images were obtained using JEM-3010 (JEOL Ltd., Tokyo, Japan). STEM images were obtained using JEM-ARM200F (JEOL Ltd., Tokyo, Japan). XPS data were obtained using SIGMA PROBE (ThermoVG, U.K). ICP-AES data were obtained using OPTIMA 8300 (PerkinElmer, Waltham, MA, USA). The machines mentioned above are installed at the National Center for Inter-University Research Facilities (NCIRF) at Seoul National University. EELS images were obtained using

Themis Z (Thermo Fisher, MA, USA) installed at the Research Institute of Advanced Materials at Seoul National University.

The powder X-ray diffraction (XRD) was performed using a D8 Advance (Bruker, Billerica, MA, USA) installed at the National Instrumentation Center for Environmental Management (NICEM) at Seoul National University. Fourier-transform infrared spectroscopy images were obtained using Spectrum Two (PerkinElmer, Waltham, MA, USA) installed at Seoul National University.

General procedure of the synthesis of amines

A glass vial (10 mL) were charged with an imine (0.20 mmol), PdCo-Fe₃O₄ (2.0 mol%), and *N,N*-dimethylacetamide (1.0 mL). The mixture was sonicated for 1 min and stirred at room temperature. Next, the vial was purged with H₂ using a balloon filled with H₂ for 1 min and stirred for 18 h at room temperature. The reaction mixture was extracted by ethyl acetate and the organic layer was filtered through a layer of Celite® and magnesium sulfate. The crude product was purified by column chromatography.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 H. Luo, Y. Yang, B. Yang, Z. Xu and D. Wang, *J. Chem. Res.*, 2021, 708–715.
- 2 (a) S. Farshbaf, L. Sreerama, T. Khodayari and E. Vessally, *Chem. Rev. Lett.*, 2018, **1**, 56–67; (b) S. Nandi, P. Patel, N.-u. H. Khan, A. V. Biradar and R. I. Kureshy, *Inorg. Chem. Front.*, 2018, **5**, 806–813.
- 3 H. M. Pinheiro, E. Touraud and O. Thomas, *Dyes Pigm.*, 2004, **61**, 121–139.
- 4 P. L. Goldschmidt, L. Savary and P. Simon, *Prog. Neuro-Psychopharmacol. Biol. Psychiatry*, 1984, **8**, 257–261.
- 5 W. J. Cubała, K. J. Wojten, A. Burkiewicz and A. Wrońska, *Psychiatr. Danubina*, 2011, **23**, 198–199.
- 6 C. M. Oliphant and G. M. Green, *Am. Fam. Physician*, 2002, **65**, 455–464.
- 7 H. d'A. Heck, S. E. Buttrill Jr, N. W. Flynn, R. L. Dyer, M. Anbar, T. Cairns, S. Dighe and B. E. Cabana, *J. Pharmacokinet. Pharmacodyn.*, 1979, **7**, 233–248.
- 8 J. K. Aronson, *Meyler's Side Effects of Drugs: The International Encyclopedia of Adverse Drug Reactions and Interactions*, Elsevier, 2015, p. 985.
- 9 (a) A. M. Tafesh and J. Weiguny, *Chem. Rev.*, 1996, **96**, 2035–2052; (b) A. Trowbridge, S. M. Walton and M. J. Gaunt, *Chem. Rev.*, 2020, **120**, 2613–2692.



- 10 (a) B. S. Takale, S. M. Tao, X. Q. Yu, X. J. Feng, T. Jin, M. Bao and Y. Yamamoto, *Org. Lett.*, 2014, **16**, 2558–2561; (b) J. B. Rivera, Y. Xu, M. Wills and V. K. Vyas, *Org. Chem. Front.*, 2020, **7**, 3312–3342.
- 11 (a) K. A. Schellenberg, *J. Org. Chem.*, 1963, **28**, 3259–3261; (b) R. P. Tripathi, S. S. Verma, J. Pandey and V. K. Tiwari, *Curr. Org. Chem.*, 2008, **12**, 1093–1115; (c) H. Alinezhad, H. Yavari and F. Salehian, *Curr. Org. Chem.*, 2015, **19**, 1021–1049.
- 12 (a) R. F. Borch, M. D. Bernstein and H. D. Durst, *J. Am. Chem. Soc.*, 1971, **93**, 2897–2904; (b) C. F. Lane, *Synthesis*, 1975, 135–146.
- 13 (a) G. W. Gribble and C. F. Nutaitis, *Org. Prep. Proced. Int.*, 1985, **17**, 317–384; (b) A. F. A. Magid, C. A. Mayanoff and K. G. Carson, *Tetrahedron Lett.*, 1990, **31**, 5595–5598; (c) A. F. A. Magid, K. G. Carson, B. D. Harris, C. A. Maryanoff and R. D. Shah, *J. Org. Chem.*, 1996, **61**, 3849–3862; (d) G. W. Gribble, *Chem. Soc. Rev.*, 1998, **27**, 395–404; (e) A. F. A. Magid and S. J. Mehrman, *Org. Process Res. Dev.*, 2006, **10**, 971–1031; (f) G. W. Gribble, *Org. Process Res. Dev.*, 2006, **10**, 1062–1075.
- 14 (a) X. Y. Zhou, M. Bao and Y. G. Zhou, *Adv. Synth. Catal.*, 2011, **353**, 84–88; (b) H. P. Hemantha and V. V. Sureshbabu, *Org. Biomol. Chem.*, 2011, **9**, 2597–2601; (c) S. M. Islam, A. S. Roy, P. Mondal and N. Salam, *Appl. Organomet. Chem.*, 2012, **26**, 625–634; (d) H. A. Patel, M. Rawat, A. L. Patel and A. V. Bedekar, *Appl. Organomet. Chem.*, 2019, **33**, e4767; (e) X. Jv, S. Sun, Q. Zhang, M. Du, L. Wang and B. Wang, *ACS Sustainable Chem. Eng.*, 2020, **8**, 1618–1626.
- 15 (a) D. Kong, M. Li, G. Zi, G. Hou and Y. He, *J. Org. Chem.*, 2016, **81**, 6640–6648; (b) H.-U. Blaser, B. Pugin, F. Spindler and A. Togni, *C. R. Chim.*, 2002, **5**, 379–385; (c) Y. Motoyama, M. Taguchi, N. Desmira, S.-H. Yoon, I. Mochida and H. Nagashima, *Chem.-Asian J.*, 2014, **9**, 71–74.
- 16 (a) D. Spasyuk, S. Smith and D. G. Gusev, *Angew. Chem., Int. Ed.*, 2013, **52**, 2538–2542; (b) M. H. Juárez, J. L. Serrano, P. Lara, J. P. M. Cerón, M. Vaquero, E. Álvarez, V. Salazar and A. Suárez, *Chem.-Eur. J.*, 2015, **21**, 7540–7555; (c) D. Deng, Y. Kita, K. Kamata and M. Hara, *ACS Sustainable Chem. Eng.*, 2019, **7**, 4692–4698.
- 17 (a) D. Wei, A. B. Voisine, D. A. Valyaev, N. Lugan and J.-B. Sortais, *Chem. Commun.*, 2018, **54**, 4302–4305; (b) U. Chakraborty, E. R. Rodriguez, S. Demeshko, F. Meyer and A. J. von Wangelin, *Angew. Chem., Int. Ed.*, 2018, **57**, 4970–4975.
- 18 (a) A. Dybov, O. Blacque and H. Berke, *Eur. J. Inorg. Chem.*, 2011, 652–659; (b) S. Chakraborty, O. Blacque, T. Fox and H. Berke, *Chem.-Asian J.*, 2014, **9**, 328–337.
- 19 (a) D. J. C. Hamilton, *Science*, 2003, **299**, 1702–1706; (b) R. H. Crabtree, *Chem. Rev.*, 2012, **112**, 1536–1554.
- 20 (a) H.-L. Jiang and Q. Xu, *J. Mater. Chem.*, 2011, **21**, 13705–13725; (b) A. K. Singh and Q. Yu, *ChemCatChem*, 2013, **5**, 652–676.
- 21 S. Byun, Y. Song and B. M. Kim, *ACS Appl. Mater. Interfaces*, 2016, **8**, 14637–14647.
- 22 J. Jang, S. Byun, B. M. Kim and S. Lee, *Chem. Commun.*, 2018, **54**, 3492–3495.
- 23 J. H. Cho, S. Byun, A. Cho and B. M. Kim, *Catal. Sci. Technol.*, 2020, **10**, 4201–4209.
- 24 A. Cho, S. Byun and B. M. Kim, *Adv. Synth. Catal.*, 2018, **360**, 1253–1261.
- 25 A. Cho, S. Byun, J. H. Cho and B. M. Kim, *ChemSusChem*, 2019, **12**, 2310–2317.
- 26 S. Yang, A. Cho, J. H. Cho and B. M. Kim, *Nanomaterials*, 2021, **11**, 2101–2115.
- 27 (a) B. S. Kumar, P. Puthiaraj, A. J. Amali and K. Pitchumani, *ACS Sustainable Chem. Eng.*, 2018, **6**, 491–500; (b) Z. Zhang, S. Liu, X. Tian, J. Wang, P. Xu, F. Xiao and S. Wang, *J. Mater. Chem. A*, 2017, **5**, 10876–10884; (c) G.-R. Xu, C.-C. Han, Y.-Y. Zhu, J.-H. Zeng, J.-X. Jiang and Y. Chen, *Adv. Mater. Interfaces*, 2018, **5**, 1701322; (d) C. H. Wu, C. Liu, D. Su, H. L. Xin, H.-T. Fang, B. Eren, S. Zhang, C. B. Murray and M. B. Salmeron, *Nat. Catal.*, 2019, **2**, 78–85; (e) H. Xue, J. Tang, H. Gong, H. Guo, X. Fan, T. Wang, J. He and Y. Yamauchi, *ACS Appl. Mater. Interfaces*, 2016, **8**, 20766–20771; (f) Y. Chen, J. Mao, R. Shen, D. Wang, Q. Peng, Z. Yu, H. Guo and W. He, *Nano Res.*, 2017, **10**, 890–896; (g) M. A. Ehsan, A. S. Hakeem and A. Rehman, *Sci. Rep.*, 2020, **10**, 14469–14479; (h) E. C. Avila, E. J. R. Ruiz, A. H. Ramirez, F. J. R. Varela, M. D. M. Acosta and D. M. Acosta, *Int. J. Hydrogen Energy*, 2017, **42**, 30349–30358.
- 28 (a) P. Yu, J. Ma, R. Zhang, J. Z. Zhang and G. G. Botte, *ACS Appl. Energy Mater.*, 2018, **1**, 267–272; (b) S. Sankar, G. M. Anilkumar, T. Tamaki and T. Yamaguchi, *ACS Appl. Energy Mater.*, 2018, **1**, 4140–4149; (c) D. Liu, Q. Guo, H. Hou, O. Niwa and T. You, *ACS Catal.*, 2014, **4**, 1825–1829; (d) C. Xu, Y. Liu, H. Zhang and H. Geng, *Chem.-Asian J.*, 2013, **8**, 2721–2728; (e) W. Ye, X. Shi, Y. Zhang, C. Hong, C. Wang, W. M. Budzianowski and D. Xue, *ACS Appl. Mater. Interfaces*, 2016, **8**, 2994–3002; (f) S. Zhang, W. Wang, Y. Gao, S. Deng, L. Ding, H. Zhuo, Z. Bao, W. Ji, C. Qiu and J. Wang, *Appl. Surf. Sci.*, 2021, **567**, 150680.
- 29 (a) T. Li, R. Wang, M. Yang, S. Zhao, Z. Li, J. Miao, Z.-D. Gao, Y. Gao and Y.-Y. Song, *Sustainable Energy Fuels*, 2020, **4**, 380–386; (b) S. Sobhani, H. Zarei and J. M. Sansano, *Sci. Rep.*, 2021, **11**, 17025–17045; (c) L. Zhang, L. Wan, Y. Ma, Y. Chen, Y. Zhou, Y. Tang and T. Lu, *Appl. Catal., B*, 2013, **138**, 229–235; (d) Y. Li, H. Zhou, T. Li, X. Jian, Z. Gao and Y.-Y. Song, *J. Mater. Chem. B*, 2021, **9**, 2016–2024; (e) Z.-Y. Liu, G.-T. Fu, L. Zhang, X.-Y. Yang, Z.-Q. Liu, D.-M. Sun, L. Xu and Y.-W. Tang, *Sci. Rep.*, 2016, **6**, 32402–32411.
- 30 (a) T. Hara, T. Kaneta, K. Mori, T. Mitsudome, T. Mizugaki, K. Ebitani and K. Kaneda, *Green Chem.*, 2007, **9**, 1246–1251; (b) C. A. Ohlin, Z. Béni, G. Laurenczy, N. Ruiz and A. M. Masdeu-Bulto, *Appl. Organomet. Chem.*, 2007, **21**, 156–160; (c) B. Y. Kara, M. Yazici, B. Kilbas and H. Goksu, *Tetrahedron*, 2016, **72**, 5898–5902; (d) W.-L. Jiang, J.-C. Shen, Z. Peng, G.-Y. Wu, G.-Q. Yin, X. Shi and H.-B. Yang, *J. Mater. Chem. A*, 2020, **8**, 12097–12105.
- 31 A. K. Chakraborti, S. Bhagat and S. Rudrawar, *Tetrahedron Lett.*, 2004, **45**, 7641–7644.

